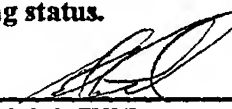


FORM PTO-1390 (REV. 11-2000)		U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NUMBER 576P043	
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371				U.S. APPLICATION NO. (If known, see 37 CFR 1.5)	
				10/018851	
INTERNATIONAL APPLICATION NO. PCT/JP00/04658		INTERNATIONAL FILING DATE 12 July 2000		PRIORITY DATE CLAIMED 14 July 1999	
TITLE OF INVENTION DYE TYPE POLARIZING PLATE					
APPLICANT(S) FOR DO/EO/US Shoji Oiso, Kumiko Ishii, Yoshitaka Kajiwara and Toru Tabei					
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:					
<p>1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.</p> <p>2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.</p> <p>3. <input checked="" type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.</p> <p>4. <input checked="" type="checkbox"/> The US has been elected by the expiration of 19 months from the priority date (Article 31).</p> <p>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2))</p> <p style="margin-left: 20px;">a. <input type="checkbox"/> is attached hereto (required only if not communicated by the International Bureau).</p> <p style="margin-left: 20px;">b. <input checked="" type="checkbox"/> has been communicated by the International Bureau.</p> <p style="margin-left: 20px;">c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US).</p> <p>6. <input checked="" type="checkbox"/> An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).</p> <p style="margin-left: 20px;">a. <input checked="" type="checkbox"/> is attached hereto.</p> <p style="margin-left: 20px;">b. <input type="checkbox"/> has been previously submitted under 35 U.S.C. 154(d)(4).</p> <p>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))</p> <p style="margin-left: 20px;">a. <input type="checkbox"/> are attached hereto (required only if not communicated by the International Bureau).</p> <p style="margin-left: 20px;">b. <input type="checkbox"/> have been communicated by the International Bureau.</p> <p style="margin-left: 20px;">c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired.</p> <p style="margin-left: 20px;">d. <input checked="" type="checkbox"/> have not been made and will not be made.</p> <p>8. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371 (c)(3)).</p> <p>9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).</p> <p>10. <input type="checkbox"/> An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</p> <p>Items 11 to 20 below concern document(s) or information included:</p> <p>11. <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.</p> <p>12. <input checked="" type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</p> <p>13. <input type="checkbox"/> A FIRST preliminary amendment.</p> <p>14. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</p> <p>15. <input type="checkbox"/> A substitute specification.</p> <p>16. <input type="checkbox"/> A change of power of attorney and/or address letter.</p> <p>17. <input type="checkbox"/> A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.</p> <p>18. <input type="checkbox"/> A second copy of the published international application under 35 U.S.C. 154(d)(4).</p> <p>19. <input type="checkbox"/> A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).</p> <p>20. <input checked="" type="checkbox"/> Other items or information:</p> <p style="margin-left: 20px;">1. Copy of the International Search Report;</p> <p style="margin-left: 20px;">2. Copy of Form PCT/IB/304; and</p> <p style="margin-left: 20px;">3. Copy of Form PCT/IB/308.</p>					

US

Annex US.II, page 2

PCT Applicant's Guide - Volume II - National Chapter - US

U.S. APPLICATION NO. 10/018851		INTERNATIONAL APPLICATION NO. PCT/JP00/04658		ATTORNEY'S DOCKET NUMBER 576P043	
21. <input checked="" type="checkbox"/> The following fees are submitted: BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)): Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO. \$1040.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$890.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$740.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$710.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00 ENTER APPROPRIATE BASIC FEE AMOUNT =				CALCULATIONS PTO USE ONLY	
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).				\$	-
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	\$	890.00
Total claims	10 - 20 =	0	x \$18.00	\$	0
Independent claims	2 - 3 =	0	x \$84.00	\$	0
MULTIPLE DEPENDENT CLAIM(S) (if applicable)				+	\$280.00
TOTAL OF ABOVE CALCULATIONS =				\$	1,170.00
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.				+	\$
SUBTOTAL =				\$	1,170.00
Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				\$	-
TOTAL NATIONAL FEE =				\$	1,170.00
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +				\$	40.00
TOTAL FEES ENCLOSED =				\$	1,210.00
				Amount to be refunded:	\$
				charged:	\$
a. <input checked="" type="checkbox"/> A check in the amount of \$ <u>1,170.00</u> & <u>40.00</u> to cover the above fees is enclosed. b. <input type="checkbox"/> Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed. c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>14-0930</u> . A duplicate copy of this sheet is enclosed. d. <input type="checkbox"/> Fees are to be charged to a credit card. WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.					
NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.					
SEND ALL CORRESPONDENCE TO: <div style="display: flex; justify-content: space-between; align-items: flex-start;"> <div style="width: 45%;"> Kevin S. Lemack Nields & Lemack 176 E. Main Street Westboro, MA 01581 </div> <div style="width: 45%; text-align: right;"> <div style="text-align: center;"></div> SIGNATURE Kevin S. Lemack NAME 32,579 REGISTRATION NUMBER </div> </div>					

DESCRIPTION

DYE TYPE POLARIZING PLATE

Technical Field

This invention relates to a novel polyvinyl alcohol type polarizing plate. Further, it relates to a polarizing plate for use in a liquid crystal projector and a color liquid crystal projector and, more in particular, it relates to a polarizing plate for use in a liquid crystal projector which is excellent both in brightness and polarizing performance and suitable to the use for a green channel, as well as a color liquid crystal projector using the polarizing plate.

Background Art

A polarizing plate having light transmission and shielding functions is a basic constituent factor, together with a liquid crystal having a light switching function, of a liquid crystal display (LCD). The application use of LCD has been extended generally from that for small sized equipments such as pocket calculators and watches at the time of initial stage to that for notebook type personal computers, word processors, liquid crystal projectors, liquid crystal television sets, car navigations and measuring instruments for indoor and outdoor use, and they are used under wide working

conditions including from low temperature to high temperature and low humidity to high humidity, so that a polarizing plate having high polarizing performance and excellent durability has been demanded.

At present, a polarizing plate is manufactured by incorporating iodine or dichroic dye as a polarizer in a stretched and oriented a film of polyvinyl alcohol or a derivative thereof. Alternatively, it is manufactured from a polyene type film by preparing a polyene through dehydrochlorination of a polyvinyl chloride film or dewatering of a polyvinyl alcohol type film and then orienting the same. Among them, the iodine type polarizing plate using iodine as a polarizer is excellent in the initial polarizing performance but is sensitive to water and heat and involves a problem in the durability when it is used under high temperature and high humidity state for a long time. For improving the durability, methods of treating with formalin or a boric acid-containing aqueous solution or using a polymer film of low moisture permeability as a protective film has been considered but this is not yet sufficient. On the other hand, a dye type polarizing plate using a dichroic dye as the polarizer is excellent in moisture proofness and heat resistance compared with the iodine type polarizing plate but the initial polarizing performance is not generally sufficient.

Further, in a polarizing plate of a neutral color prepared

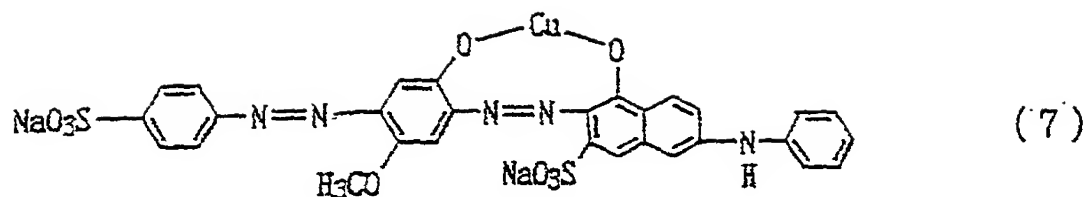
by adsorption and orientation of several kinds of dichroic dyes in a polymer film, when light leakage (color leakage) occurs at a specified wavelength in a visible light region, particularly, in a wavelength region from 400 to 700 nm in a state of stacking two polarizing plates such that their orienting directions are perpendicular to each other (crossed state), hue of the liquid crystal display sometimes changes in the dark state when the polarizing plate is attached to a liquid crystal panel. In view of the above, for preventing discoloration of the liquid crystal display due to color leakage for a specified wavelength in the dark state when the polarizing plate is attached to the liquid crystal display device, the transmittance in the crossed state (transmittance for crossed state) has to be lowered uniformly in a visible light region, particularly in a wavelength region from 400 to 700 nm.

Further, in a case of a color liquid crystal projection type display, that is, a color liquid crystal projector, a polarizing plate is used for the liquid crystal image forming portion. However, the brightness is reduced inevitably due to significant absorption of light by the polarizing plate and magnification of an image of a small area of 0.9 to 6 inch into about several tens inch to one hundred and several tens inch, so that a light source at a high luminosity is used. In addition, there is a continuous demand for the further increase of the brightness of the liquid crystal projector and, as a result,

the intensity of the light source used increases more and more.

By the way, a neutral gray iodine type polarizing plate having satisfactory polarizing performance has generally been used as a polarizing plate for a liquid crystal image forming area of a color liquid crystal projector. However, the iodine type polarizing plate involves a problem that light fastness, heat resistance and wet heat resistance are not sufficient because the polarizer is iodine. In order to solve the problem, a neutral gray polarizing plate using a dye type dichroic colorant as a polarizer has been used. In the neutral gray polarizing plate, dyes of trichromatic colors are used in combination in order to improve the transmittance and the polarizing performance in average in the entire visible ray wavelength region (400 to 700 nm). Accordingly, to the market's demand for more brightness as in the color liquid crystal projector, the light transmittance is poor and it involves a problem that the intensity of the light source has to be increased more to attain bright display. In order to solve the problem, three polarizing plates, that is, for blue channel, green channel and red channel corresponding to the trichromatic colors have become used.

As the dye used for the manufacture of a polarizing plate, Japanese Patent Publication No. 2844360 describes, for example, a water soluble dye of the following formula (7) in Example 1.



However, the polarizing plate containing the existent water soluble dye described above has not yet completely satisfied the needs of users sufficient in view of the polarization characteristics, absorption wavelength region and the hue.

Further, among the three polarizing plates corresponding to trichromatic colors of a color liquid crystal projector, that is, for blue channel, green channel and red channel, the polarizing plate for use in the green channel (green light) lacks both in favorable brightness and polarizing performance and improvement therefor has been demanded.

An object of this invention is to provide a high performance polarizing plate having excellent polarizing performance, moisture proofness and heat resistance. Further, another object of the invention is to provide a high performance neutral color polarizing plate formed by adsorbing and orienting two or more kinds of dichroic dyes in a polymer film, which causes no color leakage at the crossed state in a visible light region, particularly, in a wavelength region of from 400 to 700 nm and has excellent polarizing performance and moisture proofness and heat resistance.

A further object is to provide a high performance

polarizing plate excellent both in brightness and polarizing performance as a polarizing plate for use in green channel (green color light) among the three polarizing plates corresponding to the trichromatic colors of a colored liquid crystal projector.

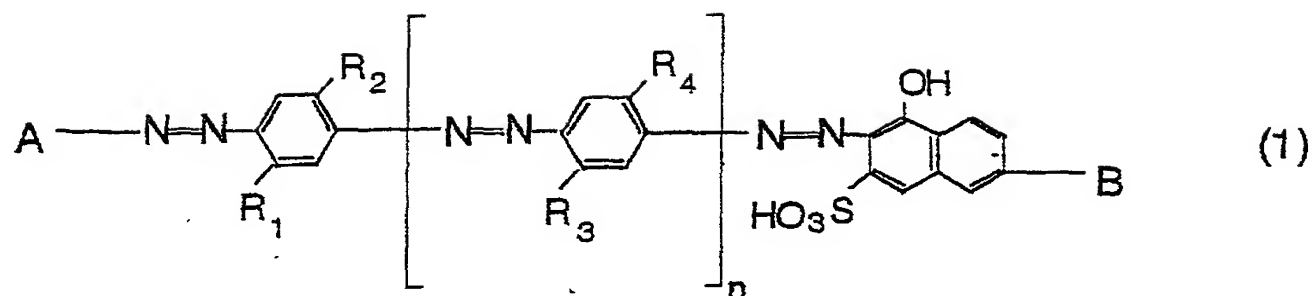
Disclosure of the Invention

In order to attain the objects described above, the present inventors have made an earnest study and, as a result, have found that a polarizing film and a polarizing plate containing a specified dye has excellent polarizing performance and moisture proofness and heat resistance, and further found that a polarizing plate excellent in the polarizing performance and the durability and with less color leakage in the visible light region can be obtained by incorporating a specified selected dye for providing a polarizing plate having a neutral color together with the above-mentioned specified dye and further that a polarizing film and a polarizing plate favorable both in the brightness and the polarizing performance suitable for use in a green channel (for green light) can be obtained by incorporating a specified dichroic dye alone or together with a selected other dye and have accomplished this invention.

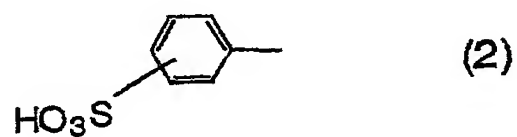
That is, this invention concerns:

1. A polyvinyl alcohol type polarizing film containing, in a substrate for the polarizing film, a water soluble dye represented by the following formula (1) in the form of a free

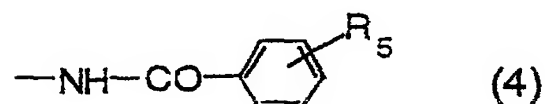
acid:



[where A represents the following formula (2)]

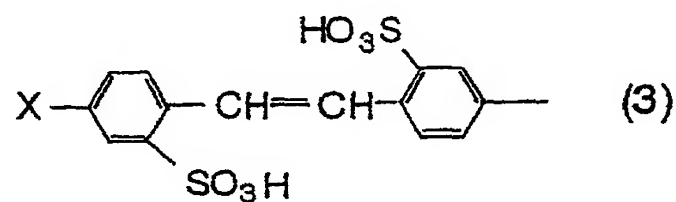


B represents the following formula (4) when A represents the formula (2);

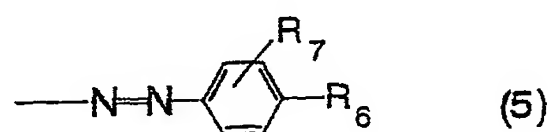


and n is 1, R₅ in the formula (4) represents an amino group or a hydroxyl group, or

A represents the following formula (3)



{where X represents a nitro group or an amino group}, B represents the following formula (5) when A represents the formula (3):

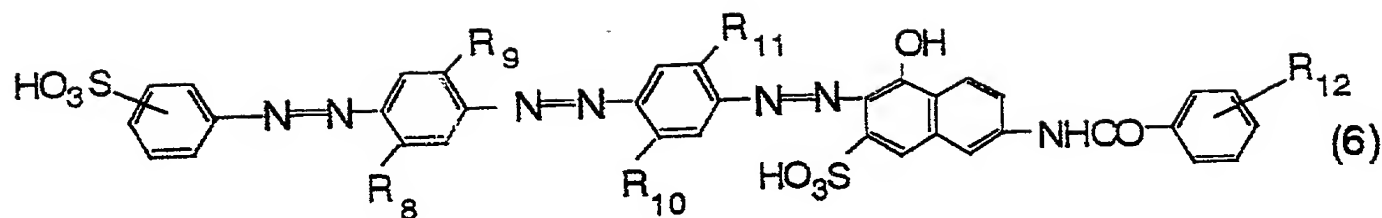


and n is 0 or 1 {where R₆ represents a hydrogen atom, hydroxyl group, substituted or unsubstituted amino group, methyl group, ethyl group, methoxy group or ethoxy group, R₇ represents a hydrogen atom, hydroxyl group, substituted or unsubstituted amino group, methyl group, ethyl group, methoxy group or ethoxy group in the formula (5)}, R₁, R₂, R₃, R₄ each independently represents a hydrogen atom, methyl group, ethyl group, methoxy group, ethoxy group and acetyl amino group, on the proviso of excluding the case where all of R₁, R₂, R₃ and R₄ are methyl group or methoxy group and a case where R₁ and R₃ are methyl group and R₂ and R₄ are methoxy group when n is 1, A represents the formula (2) and B represents the formula (4)]
or a copper complex salt dye thereof.

2. A polyvinyl alcohol type polarizing film as set forth in article 1 above, having at least one of the water soluble dyes represented by the formula (1) or the copper complex salt thereof and at least one of organic dyes other than above.

3. A polyvinyl alcohol type polarizing plate having a protective film on the surface of the polyvinyl alcohol type polarizing film as set forth in any one of articles 1 and 2 above.

4. Apolarizing film for polyvinyl alcohol for a green channel used for a liquid crystal projector containing, in a substrate for a polarizing film, a water soluble dye represented by the following formula (6) in a free acid form:



where (R_8 , R_9 , R_{10} , and R_{11} each represents independently a hydrogen atom, methyl group, ethyl group, methoxy group, ethoxy group and acetyl amino group. R_{12} represents a hydrogen atom, amino group or hydroxyl group) and having a maximum absorption wavelength (λ_{max}) of 520 nm or more and less than 580 nm.

5. A polarizing film for polyvinyl alcohol for a green channel used for a liquid crystal projector containing at least one of water soluble dyes represented by the formula (6) as set forth in article 4 above and at least one of organic dyes other than described above.

6. A polyvinyl alcohol type polarizing plate for a green channel used for a liquid crystal projector as set forth in article 4 or 5 above wherein the average light transmittance for the crossed state at 630 nm to 780 nm is 60% or more.

7. A polyvinyl alcohol type polarizing plate for a green channel used for a liquid crystal projector having a protective film on the surface of a polyvinyl alcohol type polarizing film as set forth in article 4 above.

8. A color liquid crystal projector having the polarizing plate as set forth in article 7 above in a green channel portion.

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The polarizing plate containing one or more of the water soluble dyes represented by the formula (1) or the copper complex salt dyes thereof is excellent in the polarizing performance and excellent in the moisture proofness and the heat resistance. Further, the polarizing plate containing one or more of the water soluble dyes represented by the formula (1) or copper complex salt dyes thereof and one or more of other organic dyes is excellent in any of the polarizing performance, the moisture proofness and the heat resistance and has a neutral color (gray).

Further, the polarizing plate used for the liquid crystal projector having polarizing film containing one or more of water soluble dyes represented by the formula (6) and having a maximum absorption wavelength (λ_{\max}) of 520 nm or more and less than 580 nm is favorable both in the brightness and the polarizing performance as a polarizing plate for the green channel (green light).

Best Mode for Practicing the Invention

At first, the water soluble dye represented by the formula (1) or the copper complex salt dye thereof used for the polarizing film of this invention is to be explained.

The water soluble dye represented by the formula (1) or the copper complex salt thereof used for the polarizing film of this invention includes two kinds of a dye group in which A represents the formula (2), B represents the formula (4) and

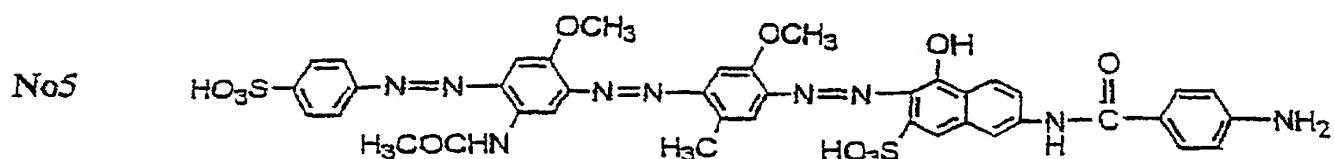
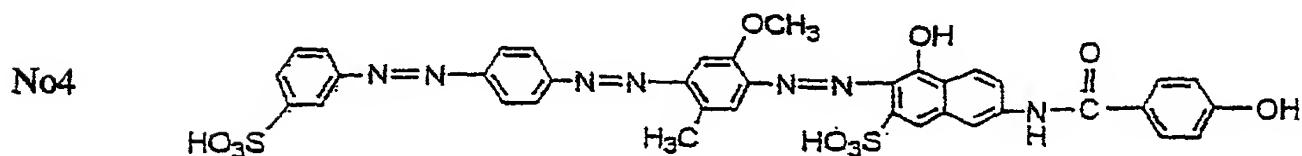
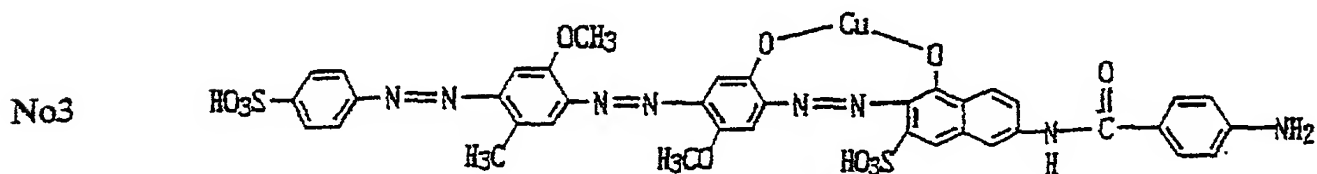
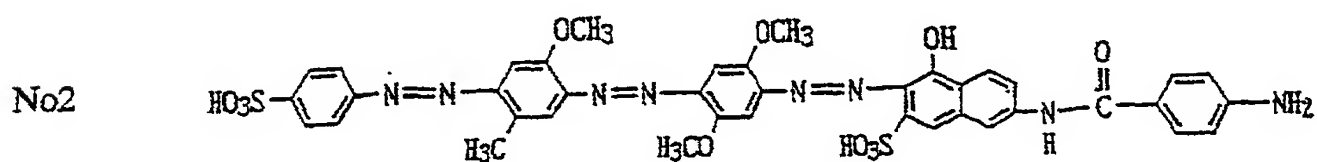
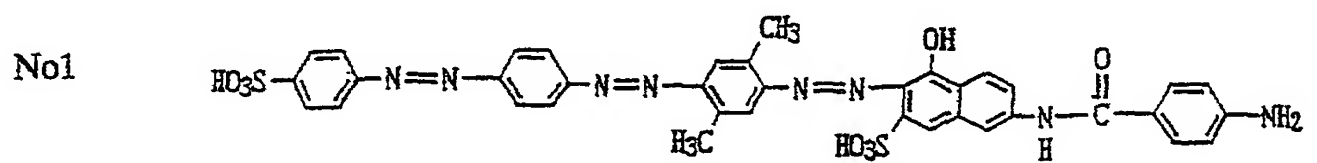
n is 1 and a dye group in which A represents the formula (3), B represents the formula (5) and n is 0 or 1, each of which may be in the form of a free acid and a salt.

The salt can include, for example, alkali metal salt such as sodium salt, potassium salt and lithium salt, ammonium salt, ethanol amine salt and alkyl amine salts, sodium salt being usually preferred.

Further, in this invention, the polarizing film containing the water soluble dye represented by the formula (1) or the copper complex salt dye thereof means a polarizing film obtained usually by dyeing the polarizing film substrate with the dye described above and stretching the same, in which the dye may be present on the surface of the film substrate or in the inside of the film, both of the cases being included.

In the dye group where A represents the formula (2), B represents the formula (4) and n is 1, among the water soluble dyes represented by the formula (1) used in this invention, the substitution position of the sulfonic group in the formula (2) is preferably the para-position relative to the azo group. R_1 , R_2 , R_3 and R_4 each represents independently a hydrogen atom, methyl group, ethyl group, methoxy group, ethoxy group or acetylamino group, and a compound in which R_1 and R_2 each represents a hydrogen atom, methyl group, methoxy group or acetyl amino group and R_3 and R_4 each represents independently a methyl group, ethyl group or methoxy group is preferred. A compound

where both of R_1 and R_2 are hydrogen atoms is more preferred. A preferred compound in this case can include, a compound where both of R_3 and R_4 are methyl group or ethyl group and a compound where both of R_3 and R_4 are methyl group is most preferred. R_5 in the formula (4) represents an amino group or hydroxyl group and the amino group is preferable. The substitution position of R_5 is particularly preferably a para-position relative to the $-NHCO-$ group. Typical examples of the dye group where A represents the formula (2), B represents the formula (4) and n is 1 are shown below in the form of free acids:

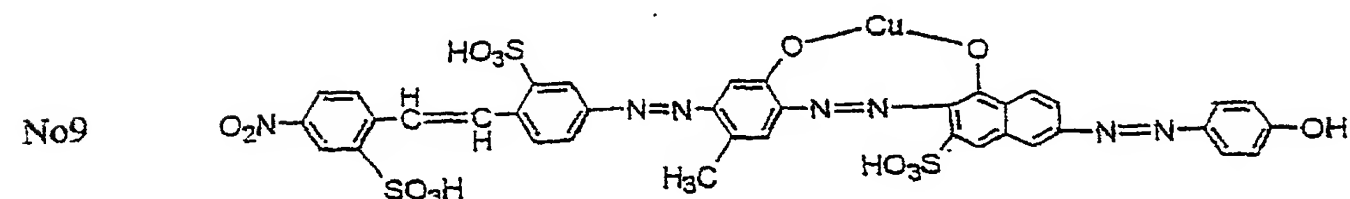
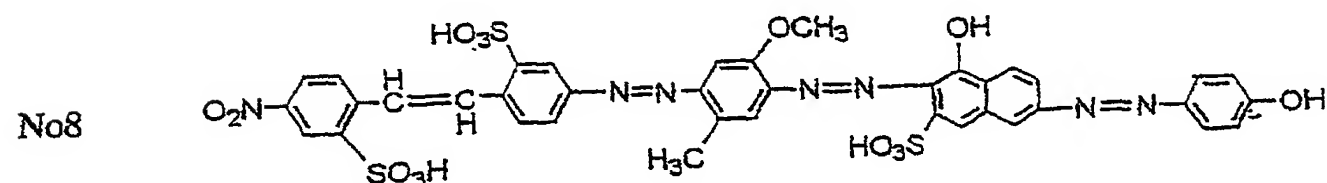
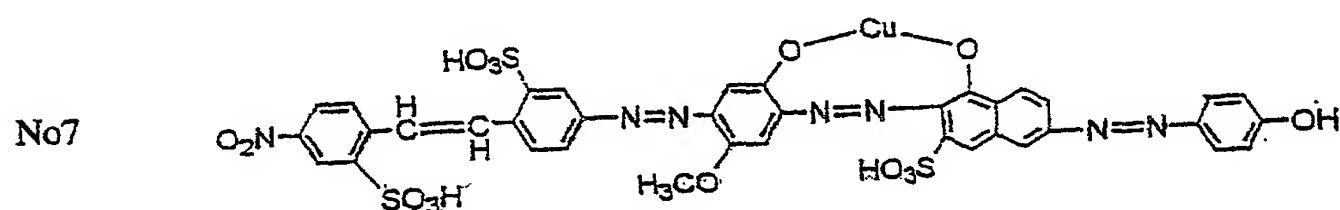
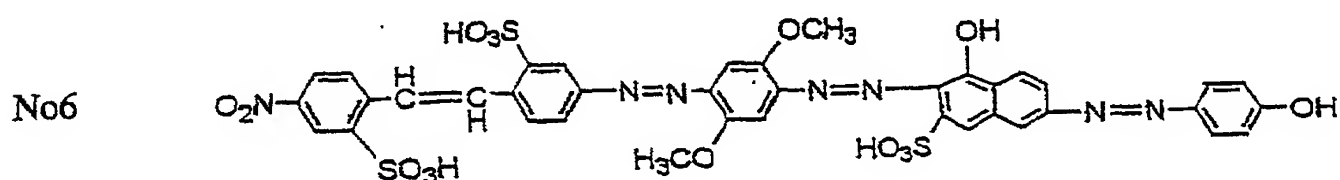


Further, in the dye group where A represents the formula (3), B represents the formula (5) and n is 0 or 1 in the formula (1) among the water soluble dyes represented by the formula (1) used in this invention, X in the formula (3) represents a nitro group or amino group, with the nitro group being preferred. R₁, R₂, R₃ and R₄ each represents independently a hydrogen atom, methyl group, ethyl group, methoxy group, ethoxy group or acetylamino group and a compounds in which R₁ and R₂ each represents a methyl group, methoxy group or acetylamino group and n is 0 is preferred and a compound in which R₁ is a methyl group or methoxy group and R₂ is a methoxy group is more preferred. Further, a compound in which both of R₁ and R₂ are methoxy group is particularly preferred. R₆ in the formula (5) represents a hydrogen atom, hydroxyl group, substituted or unsubstituted amino group, methyl group, ethyl group, methoxy group or ethoxy group, with the hydroxyl group being preferred. R₇ in the formula (5) represents a hydrogen atom, hydroxyl group, substituted or unsubstituted amino group, methyl group, ethyl group, methoxy group or ethoxy group, with the hydrogen group, hydroxyl group or carbamoyl group being preferred and the hydrogen atom or hydroxyl group being more preferred. The substituted amino group in R₆ and R₇ can include, for example, a lower alkyl amino group such as a methylamino group, a carbamoyl amino group or a lower acyl amino group such as an acetylamino group. The substitution position of R₇ is preferably the

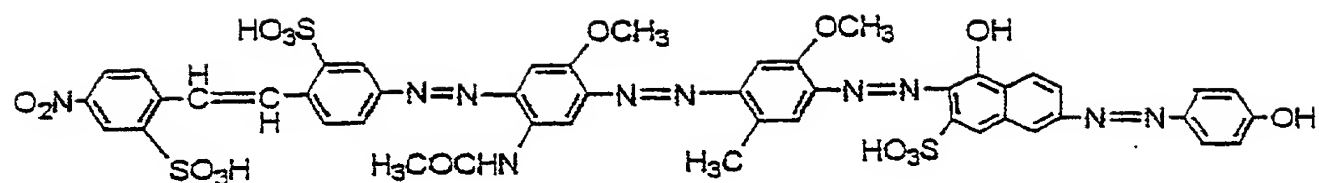
ortho-position relative to the azo group.

In this invention, the term "lower" means those having carbon atoms of from 1 to 5 and, more preferably, carbon atoms of 1 to 3.

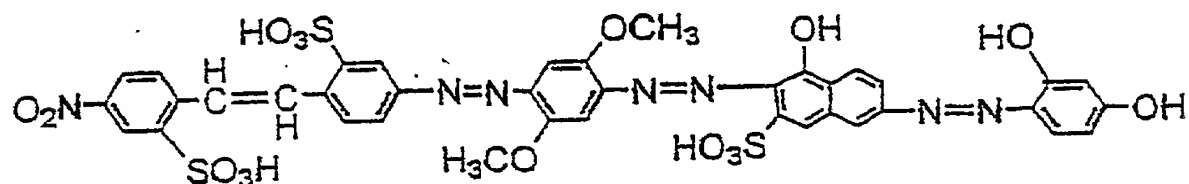
Then, among the water soluble dyes represented by the formula (1) used in this invention, typical examples of the dye group in which A represents the formula (3), B represents the formula (5) and n is 0 or 1 are shown below in the form of free acids.



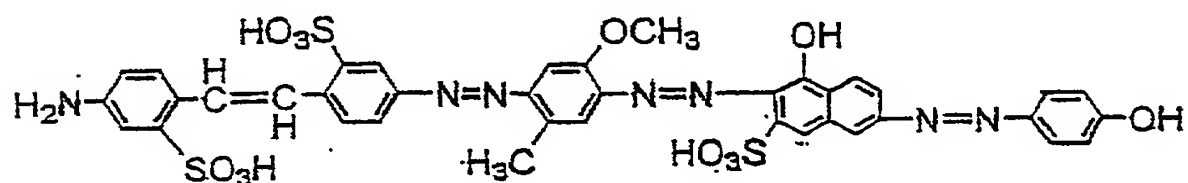
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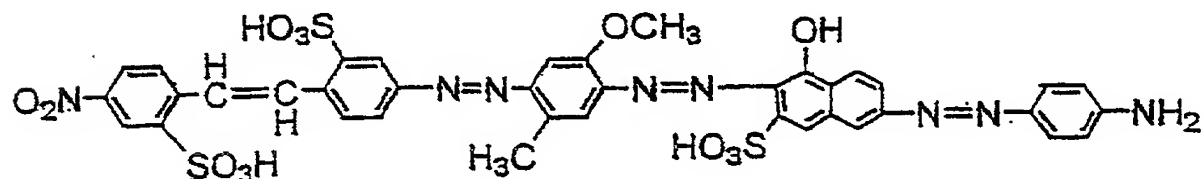
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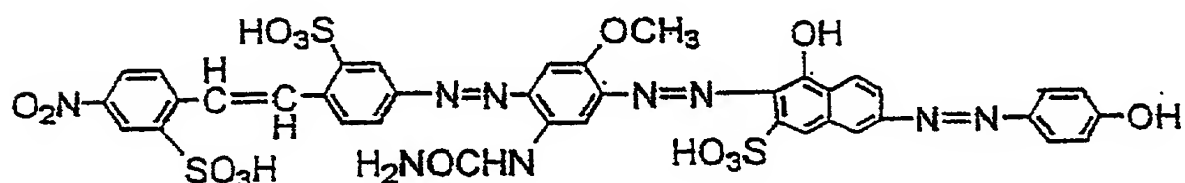
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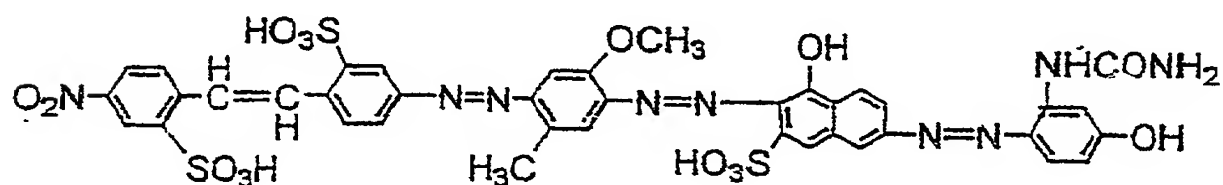
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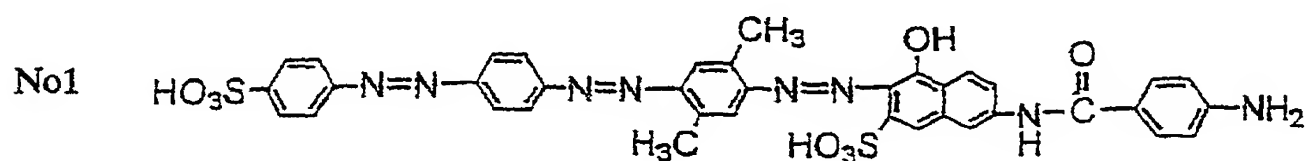


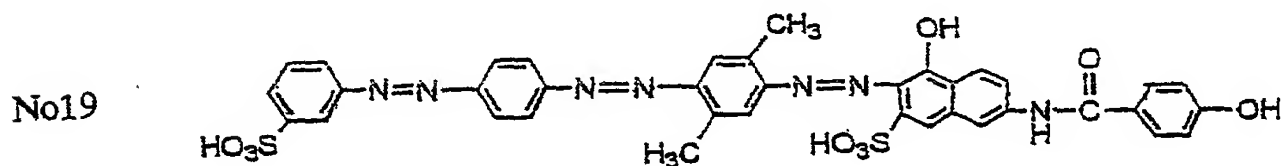
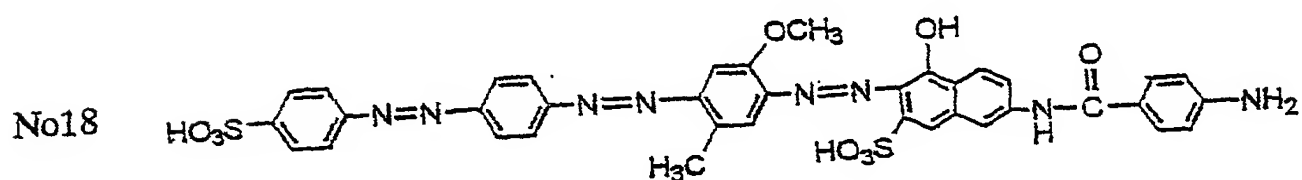
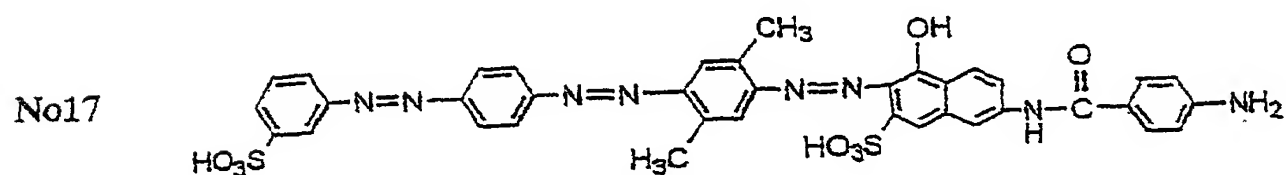
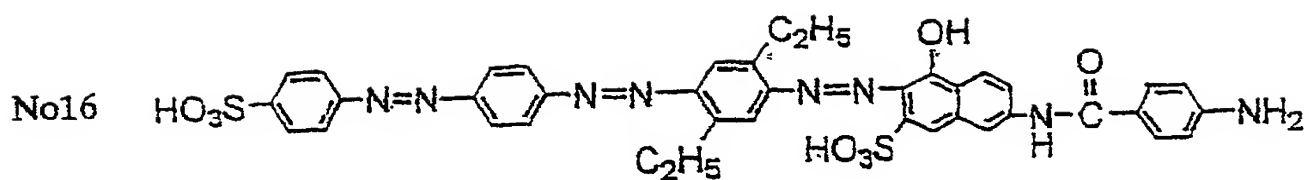
Then, description is to be made to the water soluble dye used in the polarizing plate for the green channel used for the liquid crystal projector according to this invention.

The water soluble dye used for the green channel used for the liquid crystal projector in this invention is used as a dichroic molecule and the water soluble dye can include the water soluble dye represented by the formula (6) and any of the water soluble dyes having a maximum absorption wavelength (λ_{\max}) of 520 nm or more and less than 580 nm can be used, which may be in the form of a free acid, as well as a salt. The salt can include, for example, alkali metal salts such as sodium salts, potassium salt and lithium salt, ammonium salt, ethanolamine salt and alkylamine salt, and sodium atom is preferable.

The water soluble dye represented by the formula (6) corresponds to the dye of the formula (1) in which A is a group represented by the formula (2), B is a group represented by the formula (4) and n is 1 and having a maximum absorption wavelength (λ_{\max}) of 520 nm or more and less than 580 nm and what are described for the formula (1) can be applied as they are also to the compound of the formula (6).

Then, typical examples of the water soluble dye represented by the formula (6) used in this invention are described below in the form of the free acid.





Any of the water soluble dye represented by the formula (1) used in this invention, a copper complex salt dye thereof or a water soluble dye represented by the formula (6) can be produced easily in accordance with a usual production process of azo dyes by conducting known diazotization, coupling and, optionally, formation of copper complex salt. Specific production process includes two methods for a dye group in which A represents the formula (2), B represents a formula (4) and n is 1 in the formula (1), and a dye group in which A represents the formula (3), B represents the formula (5) and n is 0 or

1. The methods are to be described more in details below.

(I) Production process for a dye group in which A represents the formula (2), B represents the formula (4) and n is 1 in the formula (1):

Anilines substituted with a sulfonic acid group of the formula (2) is diazotized, subjected to primary coupling with anilines which may have a substituent to obtain a monoazoamino compound. Then, the monoazoamino compound is diazotized and subjected to secondary coupling with aniline which may have a substituent to obtain a disazoamino compound. The disazoamino compound is diazotized and subjected to a tertiary coupling with a naphhtol corresponding to an aimed compound, that is, N-benzoyl J acid in which the phenyl nuclei of the benzoyl group is substituted with an amino or hydroxyl group in an alkaline state to obtain a water soluble dye represented by the formula (1). Further, when formation of copper complex salt is conducted by a customary method using a copper sulfate or the like, a copper complex salt of the water soluble dye represented by the formula (1) is obtained.

The starting material for the dye group in which A represents the formula (2), B represents the formula (4) and n is 1 is anilines substituted with a sulfonic acid group and the anilines can include, for example, sulfanyl acid, methanyl acid and orthanilic acid.

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The substituent of the aniline, which is a component for primary and secondary couplings and which may have a substituent, can include methyl group, ethyl group, methoxy group, ethoxy group and acetyl amino group. The substituent group may be present by one or two on the phenyl group of the aniline. The bonding position relative to the amino group is either 2-position or 3-position in a case of having one substituent and 2-position and 5-position are preferred in a case of having two substituents. The anilines which may have a substituent can include, for example, aniline, 2-methyl aniline, 3-methyl aniline, 2-ethyl aniline, 3-ethyl aniline, 2,5-dimethyl aniline, 2,5-diethyl aniline, 2-methoxy aniline, 3-methoxy aniline, 2-methoxy-5-methyl aniline, 2,5-dimethoxy aniline and 2-methoxy-5-acetyl aminoaniline. The anilines may be protected at the amino group.

The protective group can include, for example, ω -methane sulfonic acid group thereof. The aniline used for the primary coupling and the aniline used for the secondary coupling may be identical or different with each other.

The N-benzoyl J acid that is naphthols as the tertiary coupling component in which the benzoyl group is substituted with one amino group or hydroxyl group can include, N-(4'-aminobenzoyl) J acid, N-(2'-aminobenzoyl) J acid, N-(4'-hydroxybenzoyl) J acid and N-(2'-hydroxybenzoyl) J acid.

For the N-benzoyl J acids in which the benzoyl group is substituted with one amino group or hydroxy group, a compound in which R₅ is an amino group can be obtained by benzoylation of the J acid with a nitro group-substituted benzoyl chloride by a known method and then reducing the nitration product by a known method. Further, N-benzoyl J acid in which R₅ is hydroxy group can also be formed by a known method.

Further, the water soluble dye represented by the formula (6) can be produced easily by the known diazodization and coupling method in accordance with the usual production process for the azo dyes in the same manner as the production process described above, to obtain the water soluble dye of the formula (6).

(II) Production Process for dyes in which A represents the formula (3), B represents the formula (5) and n is 0 or 1.

Sodium 4-nitro-4' aminostylbene-2,2'-disulfonate in which X in the formula (3) is a nitro group is diazotized and subjected to primary coupling with anilines which may have a substituent to obtain a monoazo amino compound. Then, the monoazo amino compound is diazotized and subjected to secondary coupling with anilines which may have a substituent to obtain a disazoamino compound. The disazoamino compound is diazotized and is subjected to tertiary coupling with naphthols substituted with an azo group represented by the formula (5) in an alkaline

state to obtain a water soluble dye represented by the formula (1) where n is 1. For the water soluble dye of n is 0, when the diazotization production of the monoazoamino compound and naphthols substituted with the azo group represented by the formula (5) are subjected to a secondary coupling in an alkaline state, a water soluble dye represented by the formula (1) in which n is 0 and X is a nitro group can be obtained.

The water soluble dye represented by the formula (1) in which X is an amino group can be obtained by converting the nitro group of the water soluble dye represented by the formula (1) obtained as described above where the X is a nitro group into amino group. The nitro group can be converted into the amino group by reduction using sodium sulfide or the like.

Further, the copper complex salt of the water soluble dye represented by the formula (1) (copper complex salt dye) can be obtained by forming the copper complex salt from the water soluble dye obtained as described above by a customary method using copper sulfate or the like.

The starting material in a case where A represents the formula (3) is usually sodium 4-nitro-4'-aminostyrene 2,2'-disulfonate.

The substituent of the anilines, which may have a substituent and which are the primary or secondary coupling component, can include a methyl group, ethyl group, methoxy group, ethoxy group and acetyl amino group. The anilines may

have 1 or 2 of such substituents on the phenyl group of the anilines. The coupling position relative to the amino group is at 2-position or 3-position in a case having one substituent and 2-position and 5-position are preferred in a case of having two substituents. Anilines which may have a substituent can include, for example, aniline, 2-methylaniline, 3-methylaniline, 2-ethylaniline, 3-ethylaniline, 2,5-dimethylaniline, 2,5-diethylaniline, 2-methoxyaniline, 3-methoxyaniline, 2-methoxy-5-methylaniline, 2,5-dimethoxyaniline and 2-methoxy-5-acetylaminoaniline. The anilines may be protected at the amino group.

The protective group can include, for example, ω -methane sulfonic acid group thereof. The anilines used for the first coupling and the anilines used for the secondary coupling may be identical or different with each other.

The naphthols substituted with the azo group represented by the formula (5) used in the final coupling step in the process described above can include, for example, 6-(4'-hydroxyphenylazo)-3-sulfonic acid-1-naphthol, 6-(2'-4'-dihydroxyphenylazo)-3-sulfonic acid-1-naphthol, 6-(2'-acetylamino-4'-hydroxyphenylazo)-3-sulfonic acid-1-naphthol, 6-(4'-aminophenylazo)-3-sulfonic acid-1-naphthol and 6-(4'-N-methylaminophenylazo)-3-sulfonic acid-1-naphthol. The naphthols substituted with the azo group represented by the formula (5) can be obtained by tosylating

and diazotizing J acid by a known method, coupling with phenols and anilines and then subjecting the same to detosylation. Further, the dye group where A represents the formula (3), B represents the formula (5) and n is 0 or 1 may be formed by conducting coupling using J acid instead of using naphthols substituted with the azo group represented by the formula (5) and then converting the amino group of the J acid into the azo group of the formula (5) by a customary method. That is, it can be obtained also by diazotizing the amino group of the coupled J acid and coupling with phenol groups or anilines having the substituent represented by R_6 , R_7 .

In the production process described above, the diazotization step is conducted by a normal method of mixing a nitrite such as sodium nitrite in an aqueous solution or suspension of the diazo ingredient in a mineral acid such as hydrochloric acid or sulfuric acid, or by a backward method of adding a nitrite to a neutral or weakly alkaline aqueous solution of the diazo ingredient and mixing the same with a mineral acid. The temperature for the diazotization is appropriately from -10 to 40°C . Further, the coupling step with the anilines is conducted by mixing an aqueous acidic solution of hydrochloric acid or acetic acid with the diazo solution described above at a temperature of from -10°C to 40°C under an acidic condition of pH 2 to 7.

The monoazo compound and the disazo compound obtained

by the coupling can be put to the next step by separating it through filtration as it is or after precipitation by acidic separation or salting out, or in the form of the solution or the liquid suspension as it is. In a case where the diazonium salt is less soluble and in the form of a liquid suspension, it may be filtered to obtain a pressed cake, which can be used in the succeeding diazotization step.

The secondary or tertiary coupling reaction between the diazotization product of the monoazo amino compound or disazoamino compound and naphthols coupled with the group represented by the formula (4) or the formula (5) is conducted at a temperature of -10 to 40°C and under an alkaline condition at a pH of 7 to 10. After the completion of the reaction, the product was precipitated by salting out and recovered by filtration. Further, for the formation of a copper complex salt, an aqueous solution thereof is reacted with copper sulfate or copper chloride in the presence, for example, of ammonia, monoethanol amine or diethanol amine at 95 to 100°C, and precipitated by salting out or the like and recovered through filtration. If purification is necessary, salting out may be repeated or the product may be precipitated from water by using an organic solvent.

The organic solvent can include a water soluble organic solvent, for example, alcohols such as methanol and ethanol and ketones such as acetone.

Further, for the polarizing film or the polarizing plate of this invention, the water soluble dye represented by the formula (1) or the copper complex salt dye thereof, or the water soluble dye represented by the formula (6) may be used alone or, optionally, one or more of other organic dyes may be used together. There is no particular restriction on the organic dyes to be used and those dyes having absorption characteristics at a wavelength region different from the wavelength region of the water soluble dye or the copper complex salt dye thereof of the invention and having high dichroism are preferred. They include, for example, C.I. direct yellow 12, C.I. direct yellow 28, C.I. direct yellow 44, C.I. direct orange 26, C.I. direct orange 39, C.I. direct orange 107, C.I. direct red 2, C.I. direct red 31, C.I. direct red 79, C.I. direct red 81, C.I. direct red 247, C.I. direct green 80, C.I. direct green 59, as well as those dyes described in each of the publications of Japanese Published Unexamined Patent Application No. 145255/1984, Japanese Published Unexamined Patent Application No. 156759/1985, Japanese Published Unexamined Patent Application No. 12606/1991 and Japanese Published Unexamined Patent Application No. 218610/1999. The dyes are used as free acid or alkali metal salt, ammonium salt or salts of amines.

When other organic dyes are used together optionally, the blending ratio has no particular restriction but other organic dyes are preferably used generally within a range usually

from 0.1 to 10 parts in total based on the mass of the water soluble dye represented by the general formula (1) or the copper complex salt thereof or the water soluble dye represented by the general formula (6).

For the polarizing film of this invention or the polarizing film used for the polarizing plate for the green channel for use in the liquid crystal projector of this invention, polarizing films of various hues and neutral color can be produced by incorporating the water soluble dye represented by the formula (1), or the copper complex salt thereof, or the water soluble dye represented by the formula (6), optionally, together with other organic dyes into a polymer film as a polarizing film material by a known method. The resultant polarizing film is provided with a protective film to form a polarizing plate, optionally provided with a protective layer or an AR layer and a support, which is used for liquid crystal projectors, desk top calculators, watches, note book type personal computers, word processors, liquid crystal television sets, car navigations and instruments or displays of indoor and outdoor use.

The substrate (polymer film) used for the polarizing film of this invention is preferably a polyvinyl alcohol type substrate, and the polyvinyl alcohol type substrate can include, for example, polyvinyl alcohols or derivatives thereof, as well as modification products of one of them with olefins such as

ethylene or propylene or unsaturated carboxylic acid such as crotonic acid, acrylic acid, methacrylic acid or maleic acid. Among all, films comprising polyvinyl alcohols or derivatives thereof are used suitably with a view point of the adsorption and the orientation of the dye.

When the water soluble dye represented by the formula (1), or a copper complex salt dye thereof represented by the formula (1) and the water soluble dye represented by the formula (6) are incorporated into the polymer film described above, a method of dyeing the polymer film is usually adopted. Dyeing is conducted, for example, as described below. At first, a water soluble dye is dissolved into water to prepare a dye bath. There is no particular restriction on the concentration of the dye in the dye bath, which is usually selected within a range about from 0.001 to 10% by weight. An auxiliary agent for dyeing may be used optionally. For example, it is suitable to use sodium sulfate at a concentration of about 0.1 to 10% by weight. The polymer film is immersed in the thus prepared dye bath to dye. The dyeing temperature is preferably about from 40 to 80°C.

The dyed polymer film is stretched to orientate the water soluble dye (dichroic colorant). Any of known methods for stretching, for example, a wet method or dry method may be used. The polymer film may be stretched before dyeing depending on the case. In this case, The water soluble dye is oriented by

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dyeing. The polymer film in which the water soluble dye is incorporated and oriented is optionally applied with a post treatment such as a boric acid treatment by a known method. Such a post treatment is applied for improvement the light transmittance and the degree of polarization of the polarizing film. The condition for the boric acid treatment differs depending on the kind of the polymer film used and the kind of the dye used. The boric acid concentration in the aqueous boric acid solution is generally within a range from 0.1 to 15% by weight and, preferably from 1 to 10% by weight, and the treatment temperature is 30 to 80°C, 40 to 75°C. Further, a fixing treatment may be conducted together in an aqueous solution containing a cationic polymeric compound depending on the requirement.

A protective film having excellent optical transparency and mechanical strength can be stuck on one or both sides the thus obtained dye type polarizing film to form a polarizing plate. The material forming the protective film may be those used so far. The films include, for example, cellulose acetate type films or acrylic films, as well as fluoric type film such as of tetrafluoroethylene/ hexafluoropropylene copolymers, films comprising polyester resin, polyolefin resin or polyamide resin.

A transparent protective layer may further be provided to the surface of the polarizing plate of this invention. The

protective layer can include, for example, acrylic or polysiloxane type hard coat layer or urethane type protective layer. Further, for improving the single plate transmittance, an AR (antireflection layer) is preferably disposed on the protective layer. The AR layer can be formed, for example, by vapor depositing or sputtering of a substance such as silicon dioxide or titanium oxide and can also be formed by thinly coating a fluorine substance. An elliptic polarizing plate formed by appending a phase difference plate to the polarizing plate is also included in the polarizing plate referred to in this invention.

The thus constituted polarizing plate has a feature of having a neutral color, causing no color leakage at the crossed state in a visible light region, particularly, in a wavelength region of 400 to 700 nm, having excellent polarizing performance, causing no discoloration or lowering of the polarizing performance at higher temperature and high humidity state and with less light leakage at the crossed state in the visible light region.

The polarizing plate of this invention used for the green channel for use in the liquid crystal projector has a polarizing film containing a water soluble dye represented by the formula (6) and having a maximum absorption wavelength (λ_{\max}) at 520 nm or more and less than 580 nm as the dichroic molecule, further, optionally with other organic dyes described above. Further,

the polarizing film of this invention used in the polarizing plate for the green channel for use in the liquid crystal projector is also produced by the process described in the production process of the polarizing film according to this invention described above, further, provided with a protective film to form a polarizing plate and, optionally, provided with a protective layer, or an AR layer and a support and used as the polarizing plate used for a green channel for use in the liquid crystal projector.

The maximum absorption wavelength (λ_{max}) means the maximum absorption wavelength at the state where two polarizing plate dyed with the dichroic dye represented by the formula (6) are stacked such that the orientation directions thereof are crossed to each other.

To be used for the green channel for use in the liquid crystal projector the polarizing plate preferably has a single plate average transmittance of 39% or more and an average transmittance at the crossed state of 0.4% or less at 500 to 580 nm, and an average transmittance at the crossed state of 60% or more at 630 to 780 nm and, more preferably, the polarizing plate has a single plate average transmittance of 41% or more, and an average transmittance at the crossed state of 0.3% or less at 500 to 580 nm and, more preferably, 0.2% or less, and the average transmittance at the crossed state of 70% or more at 630 to 780 nm. Further preferably, the polarizing plate

has a single plate average transmittance of 42% or more, an average transmittance at the crossed state of 0.1% or less at 500 to 580 nm and an average transmittance at the crossed state at 630 to 780 nm of 80% or more. The polarizing plate described above exhibits red under a natural light which is called as a color polarizing plate. The color polarizing plate for the green channel of this invention has brightness and excellent polarizing performance as described above.

The polarizing plate for use in the liquid crystal projector according to this invention may be formed as a polarizing plate with an AR layer by further providing an AR layer to a polarizing plate comprising a polarizing film and a protective layer. Further, a support such as a transparent glass plate may be provided to the polarizing plate comprising the polarizing film and the protective film or the polarizing plate with the AR layer to form a polarizing plate with a support.

The single plate average transmittance is an average value of the light transmittance at a specified wavelength region upon incidence of a natural light to a polarizing plate with no AR layer and no support such as a transparent glass plate (when simply referred to as a polarizing plate, it is used hereinafter also in the same meaning). The average transmittance at the crossed state is an average value of the light transmittance at a specified wavelength region upon incidence of natural light to two polarizing plates with the

orientation directions being arranged in a crossed state.

The polarizing plate for use in the liquid crystal projector according to this invention is preferably a polarizing plate with support in which the polarizing plate is struck on the support such as a transparent glass. Further, a polarizing plate with the AR layer and the support, in which the polarizing plate having the AR layer is struck on the support, is more preferred.

The polarizing plate for use in the liquid crystal projector according to this invention is usually used as a polarizing plate with a support. The support preferably has a planar portion for sticking the polarizing plate and it is preferably a glass molded product since this is for optical application use. The glass molding product can include, for example, glass plate, lens, prism (trigonal prism, cubic prism). A lens appended with the polarizing plate can be utilized as a condenser lens with a polarizing plate in the liquid crystal projector. Further, a prism appended with the polarizing plate can be utilized as a polarizing beam splitter with a polarizing plate or a dichroic prism with a polarizing plate in the liquid crystal projector. Further, it may be appended to a liquid crystal cell. The material for the glass can include, for example, inorganic type glass such as soda glass, borosilicate glass or sapphire glass, as well as organic glass such as of acrylic or polycarbonate type. The inorganic type glass being

preferred. The thickness and the size of the glass plate may be of a desired dimension. Further, for the polarizing plate with glass, it is preferred to provide an AR layer to one or both of the surfaces of the glass surface or the polarizing plate surface in order to further improve the single plate light transmittance.

A polarizing plate with the support for use in a liquid crystal projector can be produced, for example, by coating a transparent bonding (adhesive) agent to a planar portion of a support, and then being stuck on the polarizing plate of this invention in the coated surface. Alternatively, a transparent bonding (adhesive) agent may be coated on a polarizing plate and a support may be stuck on its coated surface. The bonding (adhesive) agent used herein is, preferably, acrylic ester type. Further, when an elliptic polarizing plate is used as the polarizing plate, the surface of the plate on the side of a phase difference plate is usually stuck on the support but the side of the polarizing plate may be stuck on the glass molding product.

In the color liquid crystal projector using the polarizing plate according to this invention, the polarizing plate, more preferably, the color polarizing plate with the support described above is disposed at a green channel portion. The polarizing plate on the incident side of light is exposed to an intense light. Therefore, the temperature is elevated.

When the liquid crystal cell and the polarizing plate on the light incident side are in close contact with each other as in the usual liquid crystal display elements, heat of the polarizing plate on the light incidence side transmits to the liquid crystal cell, and the temperature of the liquid crystals in the liquid crystal cell exceeds an NI point (phase transition temperature: nematic-isotropic phase transmission point) and display is not possible. In order to avoid this, the liquid crystal cell and the polarizing plate on the light incidence side are spaced apart and air or gas is circulated by a cooling blower or the like to prevent overheating of the liquid crystal cell (which may be a water cooling system).

That is, in the color liquid crystal projector using the polarizing plate of this invention, the polarizing plate of the invention is disposed to either one or both of incidence side and emitting side of the liquid crystal cell of the green channel. The polarizing plate may or may not be in contact with the liquid crystal cell but it is preferred not to be in contact in view of the durability. In a system of using a PBS (polarizing beam splitter) behind a light source, an iodine type polarizing plate may be used or the polarizing plate of this invention may be used as the polarizing plate on the incident side. In a case where the polarizing plate is in contact with the liquid crystal cell on the emitting side, the polarizing plate of this invention having the liquid crystal cell as a

support can be used. In a case where the polarizing plate is not in contact with the liquid crystal cell, the polarizing plate of the invention using a support other than the liquid crystal cell is used preferably. Further, in view of the durability, it is preferred to dispose the polarizing plate of the invention on both of the incident side and the emitting side of the liquid crystal cell and it is further preferred to dispose the surface of the polarizing plate of this invention on the side of the liquid crystal cell and the surface of the support the side of the light source. The incident side of the liquid crystal cell means the side for the light source and the opposite side is referred to as the emitting side.

In the color liquid projector using the polarizing plate according to this invention, a UV-ray cut filter is preferably disposed between the light source and the polarizing plate with the support on the incident side. Further, the liquid crystal cell to be used is, for example, an active matrix type and is preferably formed by sealing liquid crystals between a transparent substrate formed with an electrode and TFT and a transparent substrate formed with a counter electrode. Light emitted from a light source such as a metal halide lamp is passed through the UV-ray cut filter and separated into trichromatic colors and then the green light is passed through the color polarizing plate with the support for the green channel of this invention and then joined with other dichromatic colors,

magnified by a projection lens and then projected on a screen.

The polarizing plate used for the green channel in the color liquid crystal projector thus constituted can provide a color polarizing plate for the green channel having excellent brightness and polarizing performance of 41% or more of single plate average transmittance and 0.2 or less of average transmittance at the crossed state at an absorption wavelength region of 500 to 580 nm.

Example

This invention is to be explained further specifically by way of examples but they are only illustrative and no way restrict the invention. % and parts in the examples are on the mass basis unless otherwise specified.

Synthesis Example 1, Synthesis of Compounds No. 1 and No. 16.

(a) 29.9 parts of 4-aminoazobenzene-4'-sodium sulfonate were added to 600 parts of water and dissolved at 70°C. They were cooled to 30°C or lower and 32 parts of 35% hydrochloric acid were added and then 6.9 parts of sodium nitrite were added and stirred at 25 to 28°C for 2 hours. After adding 12.1 parts of 2,5-dimethyl aniline and stirring at 25 to 30°C for 2 hours, pH was adjusted to 3 with addition of sodium carbonate and stirring was conducted further to complete coupling reaction, and filtration was conducted to obtain a disazo compound. After

dispersing the resultant disazo compound into 600 parts of water, 32 parts of 35% hydrochloric acid and then 6.9 parts of sodium nitrite were added and stirred at 25 to 30°C for 2 hours to conduct diazotize.

On the other hand, 35.8 parts of N-(4'-aminobenzoyl)J acid were added to 250 parts of water, rendered weakly alkaline with sodium carbonate and dissolved. The diazotization product of the disazo compound obtained previously was poured in this solution while being kept at neutral to weakly alkaline condition, and stirred to complete the coupling reaction. The product was salted out with sodium chloride and filtered to obtain a water soluble dye (Na salt) of trisazo compound as Compound No. 1.

(b) A water soluble dye as Compound No. 16 (Na salt) was obtained by using 2,5-diethylaniline instead of 2,5-dimethylaniline.

Synthesis Example 2, Synthesis of Compound No. 2

17.3 parts of sulfanyl acid were added to 500 parts of water and dissolved with sodium hydroxide. They were cooled to 10°C or lower and 32 parts of 35% hydrochloric acid were added and then 6.9 parts of sodium nitrite were added and stirred at 5 to 10°C for one hour. 13.7 parts of p-cresidine dissolved in an aqueous diluted hydrochloric acid were added and, while stirring at 10-15°C, adjusted to pH 3 with addition of sodium

carbonate and stirred further to complete the coupling, and filtered was conducted to obtain a monoazo compound. After dispersing the resultant monoazo compound into 600 parts of water, 32 parts of 35% hydrochloric acid and then 6.9 parts of sodium nitrite were added and stirred at 25°C to 30°C for 2 hours. 15.3 parts of 2,5-dimethoxyaniline dissolved in an aqueous diluted hydrochloric acid were added and while being stirred at 20 to 30°C, adjusted to pH to 3 with addition of sodium carbonate and stirred further to complete the coupling reaction, and filtered to obtain a disazo compound. After dispersing the resultant disazo compound into 600 parts of water, 32 parts of 35% hydrochloric acid and then 6.9 parts of sodium nitrite were added and stirred at 25 to 30°C for 2 hours to diazotize.

On the other hand, 35.8 parts of N-(4'-aminobenzoyl) J acid were added to 250 parts of water and dissolved while being rendered weakly alkaline with sodium carbonate. The diazotization product of the disazo compound obtained previously was poured into the solution while being kept at a neutral to weakly alkaline condition, and stirred to complete the coupling reaction. It was salted out by sodium chloride and filtered to obtain an aqueous dye of trisazo compound (Na salt) as Compound No. 2.

Synthesis Example 3, Synthesis of Compound No. 3

40 parts of the compound obtained in Synthesis Example 2 were dispersed in 500 parts of water and 15 parts of crystalline copper sulfate and 15 parts of monoethanol amine were added and heated to 95°C and reacted for 10 hours. The reaction solution was salted out with addition of sodium chloride and filtered to obtain a copper complex salt dye (Na salt) as Compound No. 3.

Synthesis Example 4, Synthesis of Compound No. 6

(a) 44.4 parts of sodium 4-nitro-4'-aminostylbene-2,2'-disulfonate was added to 600 parts of water and dissolved at 70°C. They were cooled to 30°C or lower and 32 parts of 35% hydrochloric acid and then 6.9 parts of sodium nitrite were added and stirred at 25 to 30°C for 2 hours. After adding 15.3 parts of 2,5-dimethoxy aniline and stirring at 25 to 30°C for 2 hours, pH was adjusted to 3 with addition of sodium carbonate and they were stirred further to complete the coupling reaction and filtered to obtain a monoazo compound. After dispersing the resultant monoazo compound into 600 parts of water, 32 parts of 35% hydrochloric acid and then 6.9 parts of sodium nitrite were added and stirred at 25 to 30°C for 2 hours to conduct diazotize and obtain a monoazo diazotization reaction solution.

On the other hand, 24.0 parts of J acid were dispersed in 250 parts of water and dissolved by adding 12 parts of sodium

hydroxide and 19 parts of tosyl chloride were added and stirred at 35 to 40°C for 1 hour. 42 parts of 35% hydrochloric acid and then 6.9 parts of sodium nitride were added to the solution and stirred at 10 to 15°C for 2 hours. Then after adding 9.4 parts of phenol, sodium hydroxide was added to adjust pH to 9 and they were stirred at 10 to 15°C to complete the coupling reaction. 6 parts of sodium hydroxide were added to the reaction solution and stirred at 80°C for 3 hours to cause hydrolysis. After cooling to 20 to 30°C, 35% hydrochloric acid was added to adjust pH to 7, then salted out with addition of sodium chloride and filtered to obtain 6-(4'-hydroxyphenylazo)-3-sulfonic acid-1-naphthol as the monoazo compound. The monoazo compound was added to 300 parts of water and dissolved with addition of sodium hydroxide.

The monoazo diazo reaction solution obtained previously was added to the solution in which the monoazo compound was dissolved while being kept at a weakly alkaline condition with sodium carbonate, and stirred at 15 to 25°C to complete the coupling reaction. The reaction solution was salted out with addition of sodium chloride and filtered to obtain a water soluble dye of trisazo compound (Na salt) as Compound No. 6.

Synthesis Example 5, Synthesis of Compound No. 7

40 parts of the compound obtained in Example 4 were dispersed in 500 parts of water, 15 parts of crystalline copper

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sulfate and 10 parts of monoethanol amine were added and heated at 95°C to react for ten hours. The reaction solution was salted out with addition of sodium chloride and filtered to obtain a water soluble dye (Na salt) as the copper complex salt dye as Compound No. 7.

Synthesis Example 6, Synthesis of Compound No. 8

Water soluble dye (Na salt) as Compound No. 8 was obtained by using p-cresidine instead of 2,5-dimethoxyaniline used in Synthesis Example 4.

Example 1

A polyvinyl alcohol of 75 μ m thickness was immersed for 4 minutes in an aqueous solution being 45°C and having the concentration of 0.03% of the dye of Compound No. 1 obtained in Synthesis Example 1 and 0.1% of sodium sulfate. The film was stretched by five times at 50°C in an aqueous 3% boric acid solution and washed with water and dried while being kept in a stretched state to obtain a polarizing film. The obtained polarizing film had a maximum absorption wavelength at 550 nm, the polarizing film had a high degree of polarization and showed durability for a long time even in a high temperature and high humidity state.

Example 2

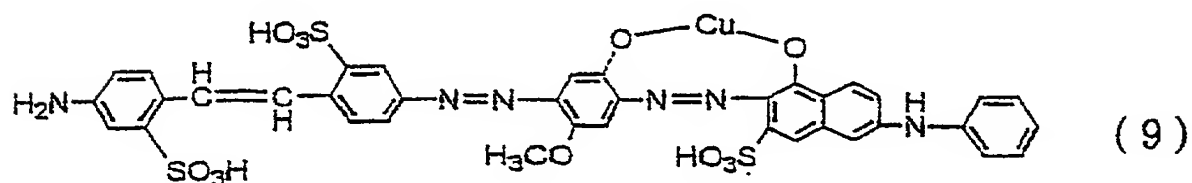
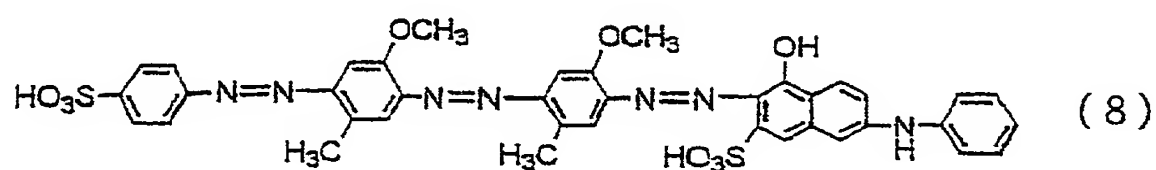
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A polyvinyl alcohol of 75 μm thickness was immersed for 4 minutes in an aqueous solution being 45°C and having the concentration of 0.03% of the dye of compound No. 3 obtained in Synthesis Example 3 and 0.1% of sodium sulfate and. The film was stretched by five times at 50°C in an aqueous 3% boric acid solution and washed with water and dried while being kept in a stretched state to obtain a polarizing film. The obtained polarizing film had a maximum absorption wavelength at 630 nm, the polarizing film had a high degree of polarization and showed durability for a long time even in a high temperature and high humidity state.

Example 3

A polyvinyl alcohol of 75 μm thickness was immersed for 4 min in an aqueous solution being 45°C and having the concentration of 0.04 % of the dye of Compound No. 1 obtained in Synthesis Example 1, 0.03% of C.I. direct orange 39, 0.04% of a dye represented by the following structural formula (8) laid-open in Example 1 of Japanese Published Unexamined Patent Application No. 12606/1991, 0.03 % of a dye represented by the following structural formula (9) laid open in Example 38 of Japanese Published Unexamined Patent Application No. 156759/1985 and 0.1% of sodium sulfate. The film was stretched by five times at 50°C in an aqueous 3% boric acid solution and washed with water and dried while being kept in a stretched

state to obtain a polarizing film. The obtained polarizing film was of a neutral color, had a high degree of polarization and showed durability for a long time even under high temperature and high humidity.



Example 4

A polyvinyl alcohol of 75 μm thickness was immersed for 4 min in an aqueous solution being 45°C having the concentration of 0.03 % of the dye of Compound No. 1, 0.04% of the dye of Compound No. 2 and 0.04% of the dye of Compound No. 3 obtained in Synthesis Examples 1 to 3, 0.03% of C.I. direct orange 39, and 0.1% of sodium sulfate. The film was stretched by five times at 50°C in an aqueous 3% boric acid solution and washed with water and dried while being kept in a stretched state to obtain a polarizing film. The obtained polarizing film was of a neutral color, had a high degree of polarization and showed durability for a long time even under high temperature and high humidity.

Example 5

A polyvinyl alcohol of 75 μm thickness was immersed for 4 minutes in an aqueous solution being 45°C and having the concentration of 0.03% of the dye of Compound No. 6 obtained in Synthesis Example 4 and 0.1% of sodium sulfate. The film was stretched by 5 times at 50°C in an aqueous 3% boric acid solution and washed with water and dried while being kept in a stretched state to obtain a polarizing film. The obtained polarizing film had a maximum absorption wavelength at 615 nm, the polarizing film had a high degree of polarization and exhibited durability for long time in a high temperature and high humidity state.

Example 6

A polyvinyl alcohol of 75 μm thickness was immersed for 4 minutes in an aqueous solution being 45°C and having the concentration of 0.03% of the dye of Compound No. 7 obtained in Synthesis Example 5 and 0.1% of sodium sulfate. The film was stretched by 5 times at 50°C in an aqueous 3% boric acid solution and washed with water and dried while being kept in a stretched state to obtain a polarizing film. The obtained polarizing film had a maximum absorption wavelength at 665 nm, the polarizing film had a high degree of polarization and exhibited durability for long time in a high temperature and

high humidity state.

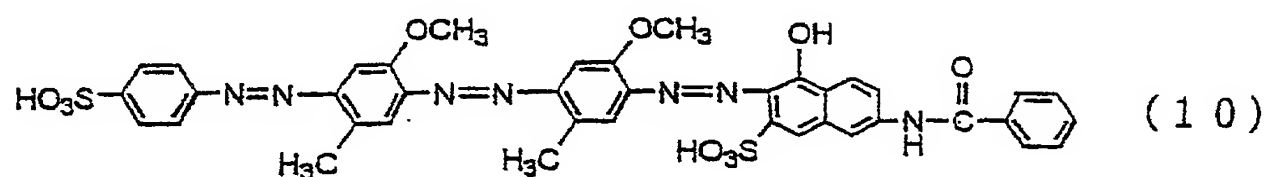
Example 7

A polyvinyl alcohol of 75 μm thickness was immersed for 4 minutes in an aqueous solution being 45°C and having the concentration of 0.05% of the dye of Compound No. 7 obtained in Synthesis Example 5, 0.04% of the dye of Compound No. 8 obtained in Synthesis Example 6, 0.03% of C.I. direct orange 39, 0.03% of C.I. direct red 81 and 0.1% of sodium sulfate. The film was stretched by five times at 50°C in a 3% aqueous boric acid solution and washed with water and dried while being kept in a stretched state to obtain a polarizing film. The obtained polarizing film exhibited a neutral color, had a high degree of polarization and exhibited durability for a long time even in a high temperature and high humid state.

Example 8

A polyvinyl alcohol of 75 μm thickness was immersed for 4 minutes in an aqueous solution being 45°C having the concentration of 0.05 % of the dye of Compound No. 7 obtained in Synthesis Example 5, 0.03% of C.I. direct orange 39, 0.03% of C.I. direct dye red 81, 0.04% of a dye shown by the following structural formula (10) as described in Example 1 of the specification of Japanese Patent Application No. 218610/1999 and 0.1% of sodium sulfate. The film was stretched by five

times at 50°C in a 3% aqueous boric acid solution and washed with water and dried while being kept in a stretched state to obtain a polarizing film. The obtained polarizing film exhibited a neutral color, had a high degree of polarization and exhibited durability for a long time even in a high temperature and high humid state.



Example 9

A polyvinyl alcohol of 75 μm thickness was immersed for 4 minutes in an aqueous solution being 45°C and having the concentration of 0.03% of the dye of Compound No. 1 obtained in Synthesis Example 1 and 0.1% of sodium sulfate. The film was stretched by five times at 50°C in an aqueous 3% boric acid solution and washed with water and dried while kept in a stretched state to obtain a polarizing film. A polarizing plate (color polarizing plate) was obtained by sticking, with a PVA type adhesive, a TAC film (80 μm thickness, trade name of products : 80 UV TAC, manufactured by Fuji Photographic Film Inc.) on one surface of the obtained polarizing film and a film prepared by forming a UV (UV-ray) curable type hard coat layer of about 10 μm thickness on one side of a TAC film on the other surface of the polarizing film. An acrylic ester type adhesive is

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applied on one side of the polarizing plate to form a color polarizing plate with an adhesive layer and, further, an AR (anti-reflection) multi-coating was applied by vacuum vapor deposition to the outside of the hard coat layer, which was cut into 30 mm to 40 mm size, and stuck on a transparent glass plate with an AR layer on one side of an identical size to obtain a color polarizing plate with AR support of the invention (for green channel of a liquid crystal projector). The color polarizing plate of this example had a maximum absorption wavelength (λ_{\max}) at 550 nm, a single plate average transmittance was 42% and an average transmittance at crossed state of 0.2% or less at 500 to 580 nm and the average transmittance at crossed state at 630 to 780 nm was 80%.

Example 10

Using the dye of Compound No. 16 obtained in Synthesis Example 1(b), a polarizing film, a color polarizing plate and a color polarizing plate with an AR support (for use in green channel of a liquid crystal projector) were prepared successively in the same method as in Example 9. The color polarizing plate of this example had a maximum absorption wavelength (λ_{\max}) at 550 nm, 42% of a single plate average transmittance, 0.2% or less of average transmittance at crossed state at 500 to 580 nm, and 80% of an average transmittance at crossed state at 630 to 780 nm.

Example 11

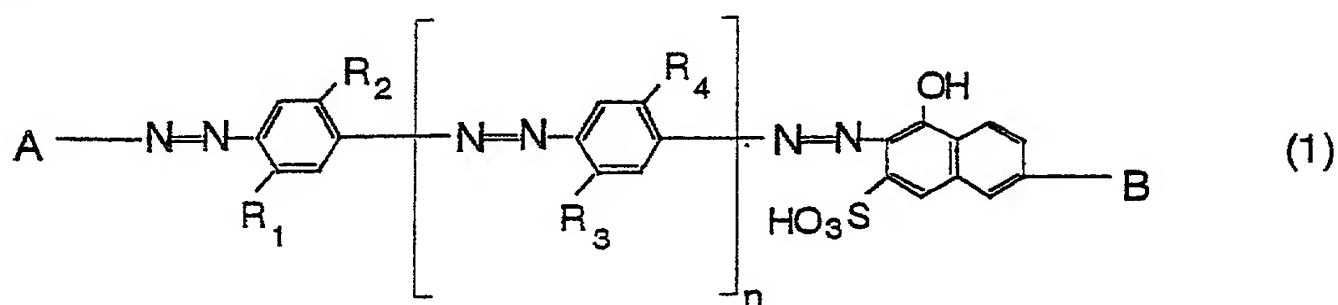
Using 0.04% of the dye of Compound No. 1 obtained in Synthesis Example 1 and 0.02% of C.I. direct orange 39, a polarizing film, a color polarizing plate and a color polarizing plate with an AR support (for green channel for use in a liquid crystal projector) were prepared successively in the same method as in Example 9. The color polarizing plate of this example had 42% of a single plate average transmittance, 0.2% or less of an average transmittance at a crossed state at 500 to 580 nm and 80% of an average transmittance at a crossed state at 630 to 780 nm.

Industrial Applicability

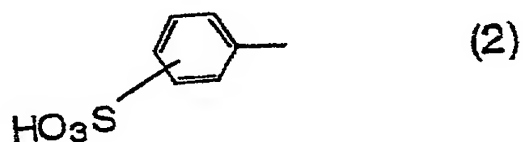
Since the polarizing plate containing the water soluble dye or the copper complex salt compound according to this invention has high polarizing performance comparable with the polarizing plate using iodine and is excellent also in durability, it is suitable to various kinds of liquid crystal displays, among all, to the display use in car-loaded application use, color polarizing plates of the green channel for use in liquid crystal projectors and industrial instruments used in various circumstances requiring high polarizing performance and durability.

Claims:

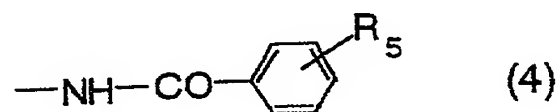
1. A polyvinyl alcohol type polarizing film containing, in a substrate for the polarizing film, a water soluble dye represented by the following formula (1) in the form of a free acid:



[where A represents the following formula (2)]

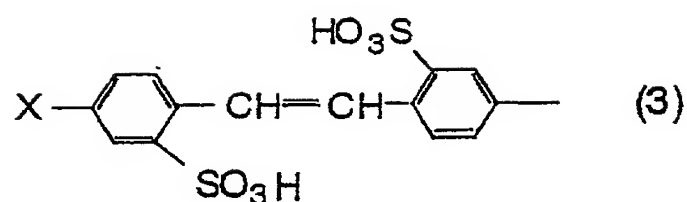


B represents the following formula (4) when A represents the formula (2);

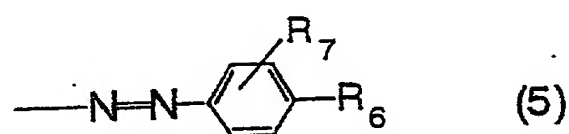


and n is 1, R₅ in the formula (4) represents an amino group or a hydroxyl group, or

A represents the following formula (3)



{where X represents a nitro group or an amino group}, B represents the following formula (5) when A represents the formula (3):



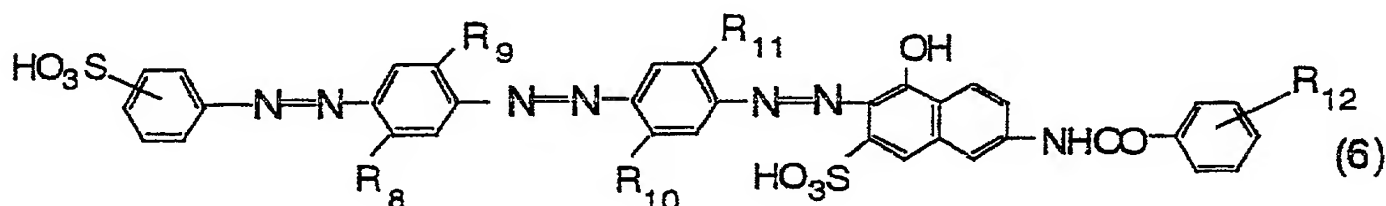
and n is 0 or 1 {where R₆ represents a hydrogen atom, hydroxyl group, substituted or unsubstituted amino group, methyl group, ethyl group, methoxy group or ethoxy group, R₇ represents a hydrogen atom, hydroxyl group, substituted or unsubstituted amino group, methyl group, ethyl group, methoxy group or ethoxy group in the formula (5)}, R₁, R₂, R₃, R₄ each independently represents a hydrogen atom, methyl group, ethyl group, methoxy group, ethoxy group and acetyl amino group, on the proviso of excluding the case where all of R₁, R₂, R₃ and R₄ are methyl group or methoxy group and the case where R₁ and R₃ are methyl group and R₂ and R₄ are methoxy group when n is 1, A represents the formula (2) and B represents the formula (4)] or a copper complex salt dye thereof.

2. A polyvinyl alcohol type polarizing film as defined in claim 1, containing at least one of the water soluble dye

represented by the formula (1) or the copper complex salt thereof and at least one of organic dyes other than above.

3. A polyvinyl alcohol type polarizing plate having a protective film on the surface of the polyvinyl alcohol type polarizing film as defined in any one of claim 1 or 2.

4. Apolarizing film for polyvinyl alcohol for a green channel used for a liquid crystal projector containing in a substrate for a polarizing film, a water soluble dye represented by the following formula (6):



where (R_8 , R_9 , R_{10} , and R_{11} each represents a hydrogen atom, methyl group, ethyl group, methoxy group, ethoxy group and acetyl amino group. R_{12} represents a hydrogen atom, amino group or hydroxyl group) in a free acid form and having a maximum absorption wavelength (λ_{max}) of 520 nm or more and less than 580 nm.

5. Apolarizing film for polyvinyl alcohol for a green channel used for a liquid crystal projector containing at least one of water soluble dyes represented by the formula (6) as defined in claim 4 and at least one of organic dyes other than described

above.

6. A polarizing plate for polyvinyl alcohol for a green channel used for a liquid crystal projector as defined in claim 4 or 5 wherein the average light transmittance for the crossed state at 630 nm to 780 nm is 60% or more.

7. A polarizing plate for polyvinyl alcohol for a green channel used for a liquid crystal projector having a protective film on the surface of a polyvinyl alcohol type polarizing film as defined in claim 4.

8. A color liquid crystal projector having the polarizing plate as defined in claim 7 in a green channel portion.

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Additional foreign applications:

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached?	
				YES	NO
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Additional provisional applications:

Application Number	Filing Date (MM/DD/YYYY)

Additional U.S. applications:

U.S. Parent Application Number	PCT Parent Number	Parent Filing Date (MM/DD/YYYY)	Parent Patent Number (if applicable)

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DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (37 CFR 1.63) <input checked="" type="checkbox"/> Declaration Submitted with Initial Filing OR <input type="checkbox"/> Declaration Submitted after Initial Filing (surcharge (37 CFR 1.16 (e)) required)	Attorney Docket Number	576P043
	First Named Inventor	Shoji OISO
	COMPLETE IF KNOWN	
	Application Number	/
	Filing Date	
	Group Art Unit	
	Examiner Name	

As a below named inventor, I hereby declare that:

My residence, mailing address, and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

DYE TYPE POLARIZING PLATE

(Title of the Invention)

the specification of which

☐ is attached hereto

OR

☐ was filed on (MM/DD/YYYY)

July 12, 2000

as United States Application Number or PCT International

(if applicable).

Application Number

PCT/JP00/04658

and was amended on (MM/DD/YYYY)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment specifically referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56, including for continuation-in-part applications, material information which became available between the filing date of the prior application and the national or PCT international filing date of the continuation-in-part application.

I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Foreign Filing Date (MM/DD/YYYY)	Priority Not Claimed	Certified Copy Attached?	
				YES	NO
11/200435	JAPAN	07/14/1999	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
11/211148	JAPAN	07/26/1999	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
11/234058	JAPAN	08/20/1999	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

☐ Additional foreign application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto:

I hereby claim the benefit under 35 U.S.C. 119(e) of any United States provisional application(s) listed below.

Application Number(s)	Filing Date (MM/DD/YYYY)	<input type="checkbox"/> Additional provisional application numbers are listed on a supplemental priority data sheet PTO/SB/02B attached hereto.

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DECLARATION — Utility or Design Patent Application

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Correspondence address below

Name Kevin S. LemackAddress Nields & LemackAddress 176 E. Main Street - Suite 8City WestboroState MAZIP 01581Country U.S.A.Telephone (508) 898-1818Fax (508) 898-2020

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C. 1001 and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

NAME OF SOLE OR FIRST INVENTOR:

☐ A petition has been filed for this unsigned inventor

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or SurnameOISOInventor's
SignatureShoji OisoDate November 6, 2001

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City

Saitama


State

ZIP

350-1108

Country

JAPAN☐ Additional inventors are being named on the _____ supplemental Additional Inventor(s) sheet(s) PTO/SB/02A attached hereto.

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ADDITIONAL INVENTOR(S)
Supplemental Sheet
Page 1 of 1

Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor					
Given Name (first and middle [if any])				Family Name or Surname			
Yoshitaka				KAJIWARA			
Inventor's Signature	Yoshitaka Kajiwarara					Date	November 6, 2001
Residence: City	Saitama	State		Country	JAPAN	Citizenship	JP
Post Office Address	6-8-25-302, Kamiochiai, Yono-shi,						
Post Office Address	Saitama 338-0001 JAPAN						
City	Saitama	State		ZIP	338-0001	Country	JAPAN
Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor					
Given Name (first and middle [if any])				Family Name or Surname			
Toru				TABELI			
Inventor's Signature	Toru Tabari					Date	November 6, 2001
Residence: City	Saitama	State		Country	JAPAN	Citizenship	JP
Post Office Address	8-90-50, Azuma, Kitamoto-shi,						
Post Office Address	Saitama 364-0007 JAPAN						
City	Saitama	State		ZIP	364-0007	Country	JAPAN
Name of Additional Joint Inventor, if any:		<input type="checkbox"/> A petition has been filed for this unsigned inventor					
Given Name (first and middle [if any])				Family Name or Surname			
Inventor's Signature						Date	
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